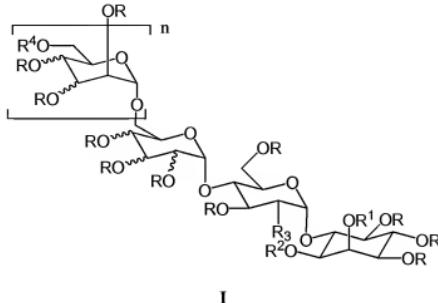


In the Claims:

1. (previously presented) A compound represented by formula I:



I

wherein,

n is 1, 3, or 4;

R represents independently for each occurrence H, alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, or -Si(alkyl);

R<sup>1</sup> and R<sup>2</sup> are independently H, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, -Si(alkyl); or R<sup>1</sup> and R<sup>2</sup> taken together are C(CH<sub>3</sub>)<sub>2</sub>, P(O)OH, or P(O)OR<sup>5</sup>;

R<sup>3</sup> is amino, -N<sub>3</sub>, or -NH<sub>3</sub>X;

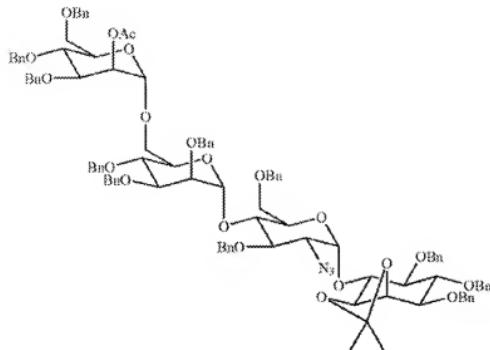
R<sup>4</sup> represents independently for each occurrence alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, -Si(alkyl)<sub>3</sub>, or -P(O)(OR<sup>5</sup>)<sub>2</sub>;

R<sup>5</sup> represents independently for each occurrence H, Li<sup>+</sup>, Na<sup>+</sup>, K<sup>+</sup>, Rb<sup>+</sup>, Cs<sup>+</sup>, aryl, or an optionally substituted alkyl group; and

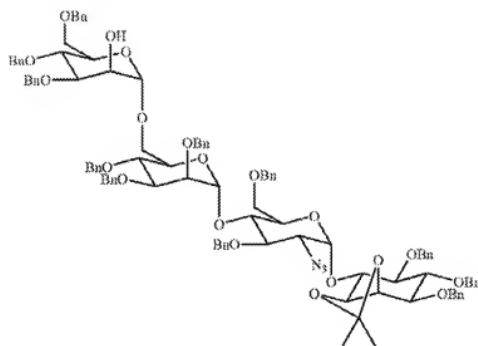
X is a halogen, alkyl carboxylate, or aryl carboxylate.

2. (canceled)
3. (original) The compound of claim 1, wherein n is 3.
4. (original) The compound of claim 1, wherein R is H.

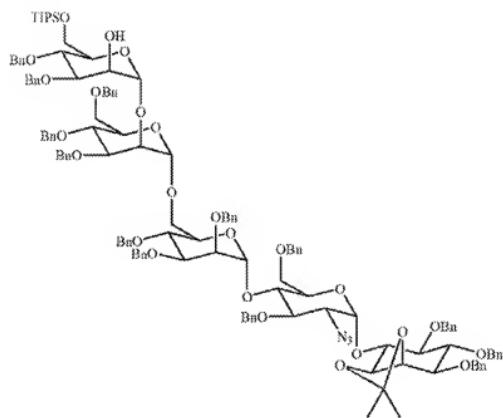
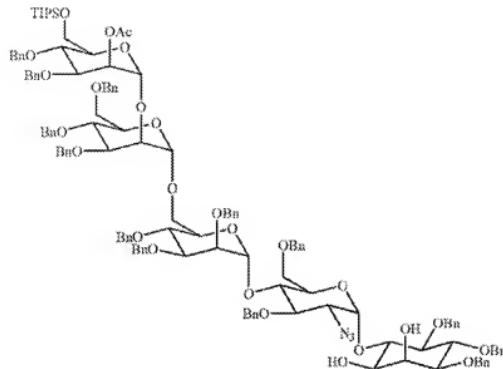
5. **(original)** The compound of claim 1, wherein R<sup>1</sup> and R<sup>2</sup> taken together are P(O)OR<sup>5</sup>.
6. **(original)** The compound of claim 1, wherein R<sup>3</sup> is N<sub>3</sub>.
7. **(original)** The compound of claim 1, wherein R<sup>3</sup> is -NH<sub>2</sub>X.
8. **(previously presented)** The compound of claim 1, wherein R<sup>4</sup> represents independently for each occurrence -CH<sub>2</sub>Ph, or -Si(alkyl)<sub>3</sub>.
9. **(previously presented)** The compound of claim 1, wherein R<sup>4</sup> represents independently for each occurrence -CH<sub>2</sub>Ph, -or P(O)OR<sup>5</sup>; and R<sup>5</sup> is an optionally substituted alkyl group.
10. **(currently amended)** A compound selected from the group consisting of:

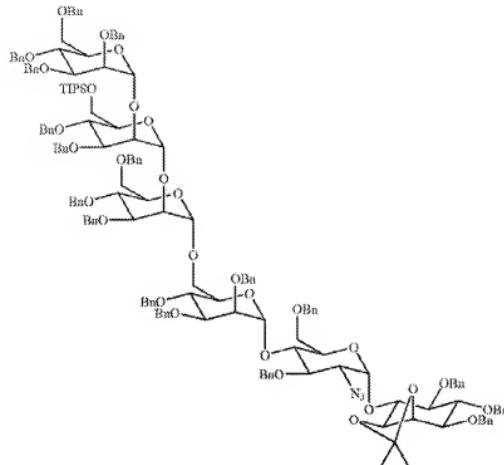


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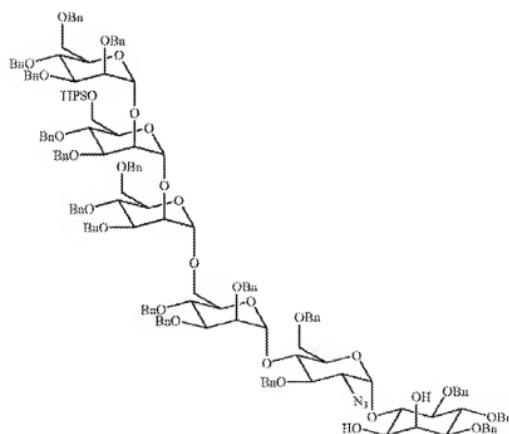


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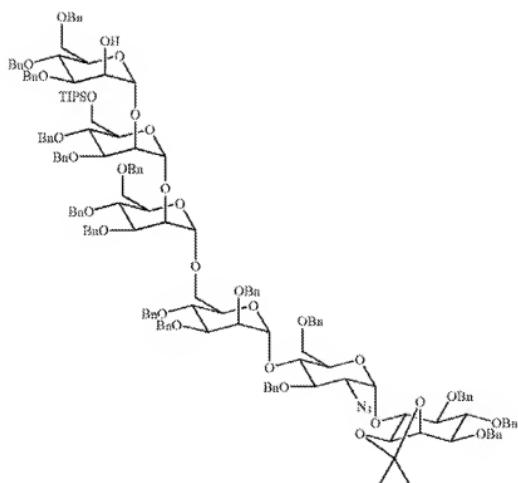
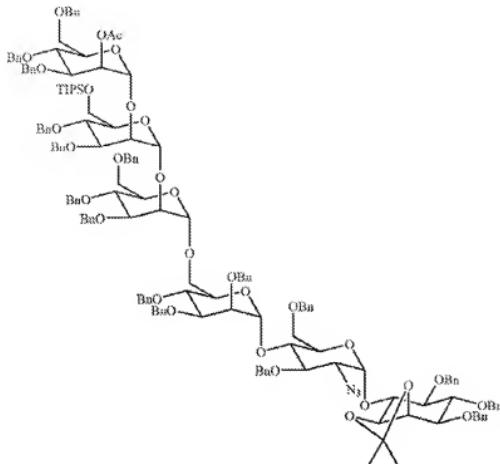


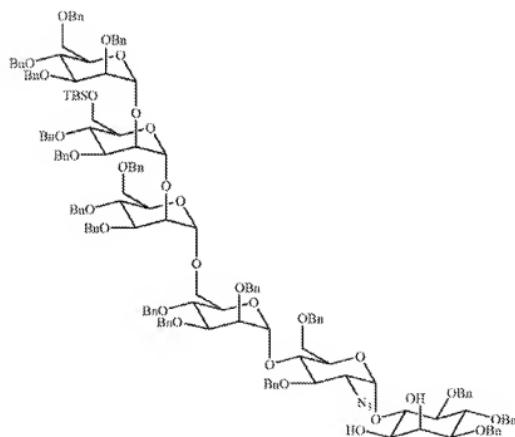
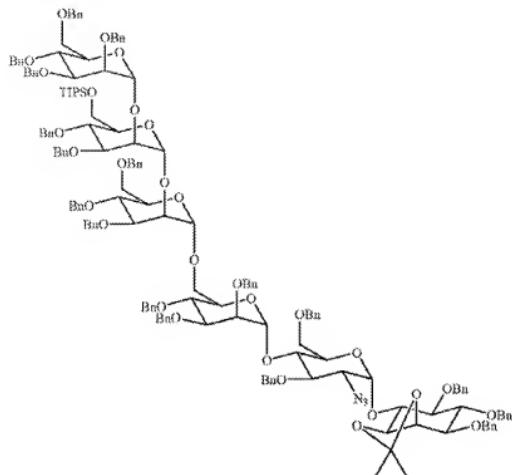


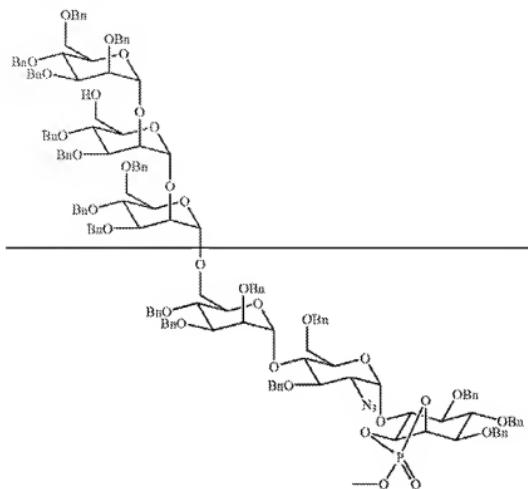
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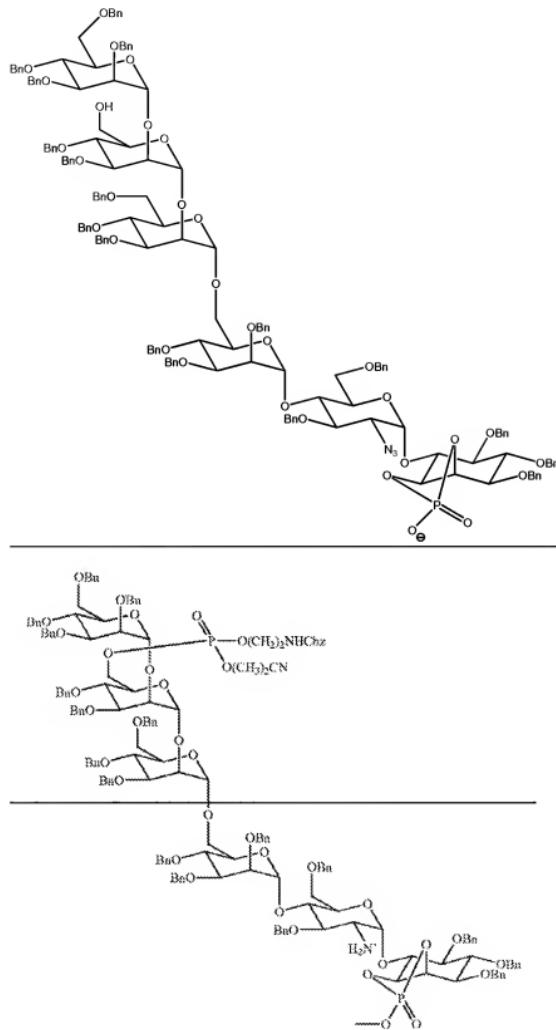


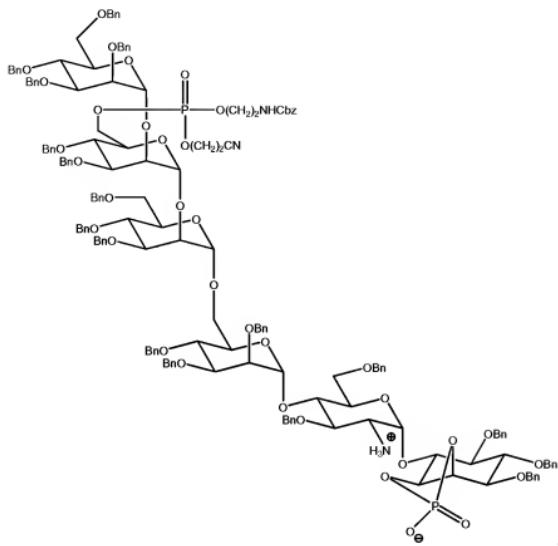
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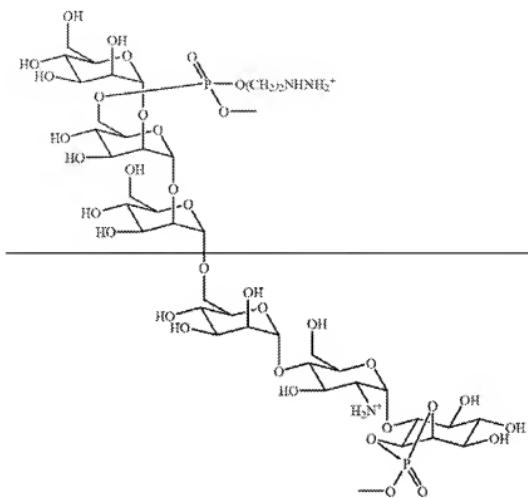


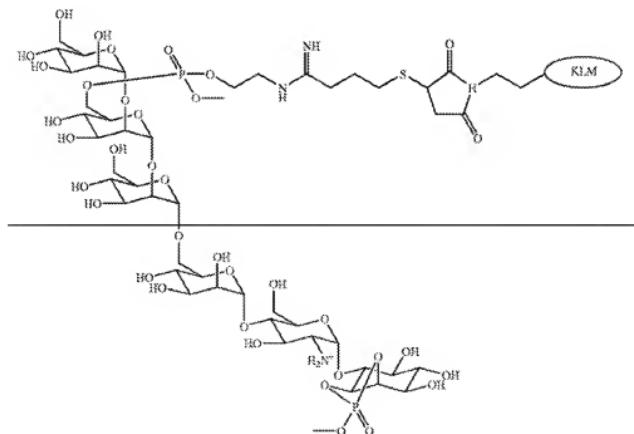
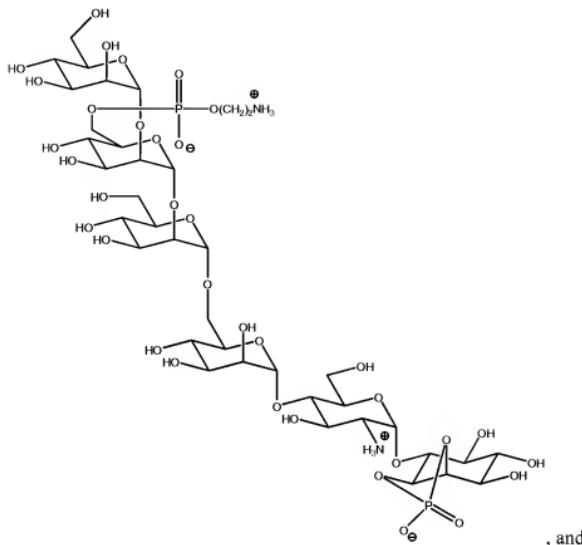


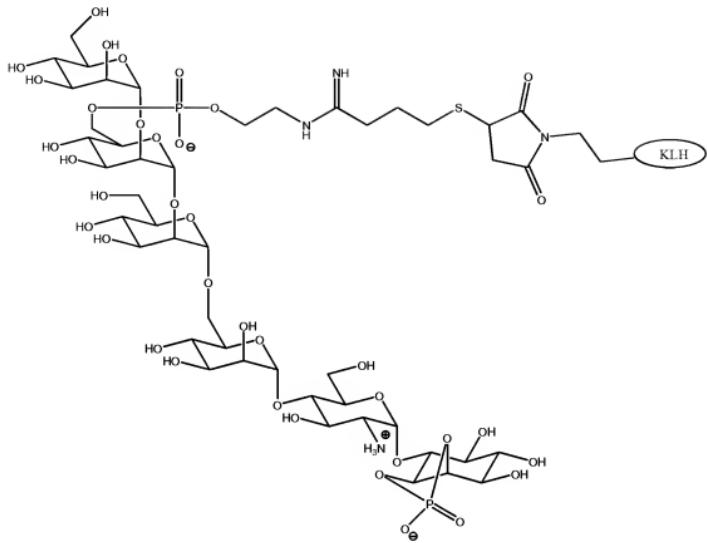




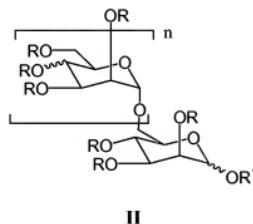








11. (previously presented) A compound represented by formula II:



II

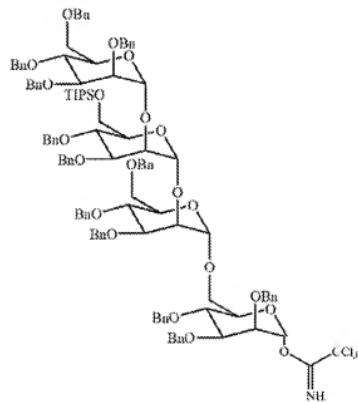
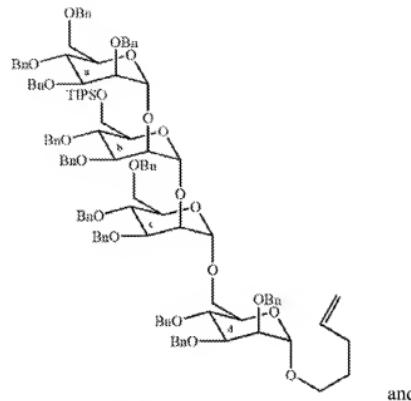
wherein,

n is 3, or 4;

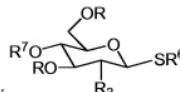
R represents independently for each occurrence H, alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, or -Si(alkyl)<sub>3</sub>;

R<sup>1</sup> is -(CH<sub>2</sub>)<sub>m</sub>CH=CH<sub>2</sub> or trichloroacetylimate; and

- m is 1-6.
12. **(canceled)**
13. **(original)** The compound of claim 11, wherein n is 3.
14. **(original)** The compound of claim 11, wherein m is 3.
15. **(original)** The compound of claim 11, wherein R represents independently for each occurrence -CH<sub>2</sub>-aryl or -Si(alkyl)<sub>3</sub>.
16. **(original)** The compound of claim 11, wherein R represents independently for each occurrence benzyl or -Si(iPr)<sub>3</sub>.
17. **(previously presented)** The compound of claim 11, wherein R<sup>1</sup> is trichloroacetimidate and R represents independently for each occurrence benzyl or -Si(iPr)<sub>3</sub>.
18. **(previously presented)** The compound of claim 11, wherein said compound of formula II is selected from the group consisting of:



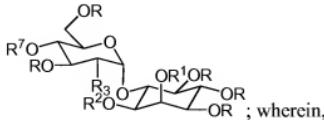
19. (previously presented) A method comprising the step of:



admixing a compound represented by with a compound represented

by , followed by the addition, together or separately of *N*-

iodosuccinimide and silver triflate, thereby forming a compound represented by



R represents independently for each occurrence H, alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, or -Si(alkyl)<sub>3</sub>;

R<sup>1</sup> and R<sup>2</sup> are independently H, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, -Si(alkyl)<sub>3</sub>; or R<sup>1</sup> and R<sup>2</sup> taken together are C(CH<sub>3</sub>)<sub>2</sub>, P(O)OH, or P(O)OR<sup>5</sup>;

R<sup>3</sup> is amino, -N<sub>3</sub>, or -NH<sub>3</sub>X;

R<sup>5</sup> represents independently for each occurrence H, Li<sup>+</sup>, Na<sup>+</sup>, K<sup>+</sup>, Rb<sup>+</sup>, Cs<sup>+</sup>, aryl, or an optionally substituted alkyl group;

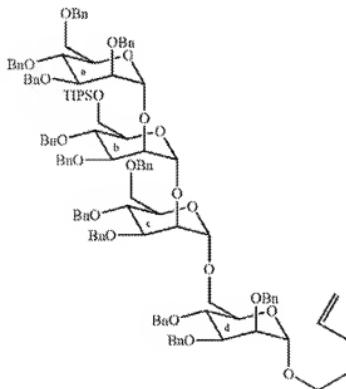
R<sup>6</sup> is alkyl or aryl;

R<sup>7</sup> is alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, or -Si(alkyl)<sub>3</sub>; and

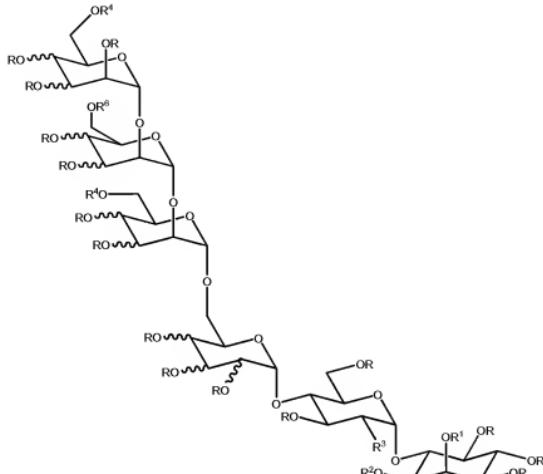
X is a halogen, alkyl carboxylate, or aryl carboxylate.

20. (original) The method of claim 19, wherein R is -CH<sub>2</sub>-aryl.
21. (original) The method of claim 19, wherein R<sup>1</sup> and R<sup>2</sup> taken together are C(CH<sub>3</sub>)<sub>2</sub>.
22. (original) The method of claim 19, wherein R<sup>3</sup> is -N<sub>3</sub>.
23. (original) The method of claim 19, wherein R<sup>6</sup> is alkyl.
24. (original) The method of claim 19, wherein R<sup>7</sup> is -C(O)-alkyl.
25. (original) The method of claim 19, wherein R is benzyl, R<sup>1</sup> and R<sup>2</sup> taken together are C(CH<sub>3</sub>)<sub>2</sub>, and R<sup>3</sup> is -N<sub>3</sub>.
26. (original) The method of claim 19, wherein R is benzyl, R<sup>1</sup> and R<sup>2</sup> taken together are C(CH<sub>3</sub>)<sub>2</sub>, R<sup>3</sup> is -N<sub>3</sub>, and R<sup>6</sup> is ethyl.
27. (previously presented) A method of preparing a tetrasaccharide, comprising the steps of:

- covalently binding a mannopyranoside to a solid support to provide a first substrate, reacting said first substrate with a mannopyranose trichloroacetimidate to give a disaccharide bound to said solid support, reacting said disaccharide with a mannopyranose trichloroacetimidate to give a triisaccharide bound to said solid support, reacting said triisaccharide with a mannopyranose trichloroacetimidate to give a tetrasaccharide bound to said solid support, and cleaving said tetrasaccharide from said solid support.
28. **(original)** The method of claim 27, wherein said mannopyranoside is bound to said solid support through a glycosidic linkage.
29. **(original)** The method of claim 27, wherein said tetrasaccharide is cleaved from said solid support using Grubbs' catalyst.
30. **(previously presented)** The method of claim 27, wherein said tetrasaccharide is



31. (new) A compound represented by formula I:



wherein,

R represents independently for each occurrence H, alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, or -Si(alkyl)<sub>3</sub>;

R<sup>1</sup> and R<sup>2</sup> are independently H, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, -Si(alkyl)<sub>3</sub>; or R<sup>1</sup> and R<sup>2</sup> taken together are C(CH<sub>3</sub>)<sub>2</sub>, P(O)OH, or P(O)OR<sup>5</sup>;

R<sup>3</sup> is amino, -N<sub>3</sub>, or -NH<sub>3</sub>X;

R<sup>4</sup> represents independently for each occurrence H, alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, -Si(alkyl)<sub>3</sub>, or -P(O)(OR<sup>5</sup>)<sub>2</sub>;

R<sup>5</sup> represents independently for each occurrence H, Li<sup>+</sup>, Na<sup>+</sup>, K<sup>+</sup>, Rb<sup>+</sup>, Cs<sup>+</sup>, aryl, or an optionally substituted alkyl group; and

R<sup>6</sup> represents independently for each occurrence alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, -Si(alkyl)<sub>3</sub>, or -P(O)(OR<sup>5</sup>)<sub>2</sub>;

- X is a halogen, alkyl carboxylate, or aryl carboxylate.
- 32. (new) The compound of claim 31, wherein R is H.
  - 33. (new) The compound of claim 31, wherein R<sup>1</sup> and R<sup>2</sup> taken together are P(O)OR<sup>5</sup>.
  - 34. (new) The compound of claim 31, wherein R<sup>3</sup> is -NH<sub>3</sub>X.
  - 35. (new) The compound of claim 31, wherein R<sup>4</sup> is H.
  - 36. (new) The compound of claim 31, wherein R<sup>6</sup> is -P(O)(OR<sup>5</sup>)<sub>2</sub>.
  - 37. (new) The compound of claim 31, wherein R is H; R<sup>1</sup> and R<sup>2</sup> taken together are P(O)OR<sup>5</sup>; R<sup>3</sup> is -NH<sub>3</sub>X; R<sup>4</sup> is H; and R<sup>6</sup> is -P(O)(OR<sup>5</sup>)<sub>2</sub>.